wherein R represents a carboxyl group or the functional derivative thereof selected from the group consisting of a carboxylic lower alkyl ester carboxylic acid aralkyl ester , and cya<u>no groups</u>; R² represents a hydrogen atom, a lower alkyl group, a lower alkoxy group, R4S(O) group [(] wherein R4 represents a lower alkyl group and n represents 0, 1 or 2[)], a lower/alkanoyl group, an aryl group, an aroyl group, a carboxyl group or the functional derivative thereof selected from the group consisting of a carboxylic acid carboxylic acid aralkyl ester radical cyano groups, a lower alkenyl group, a sulfamoyl group, ϕ r a heterocyclic residue; and R^3 represents a lower alkyl-subs∲ituted tetrazolyl group or a lower alkyl-substituted thiadiazolyl group and the pharmaceutically acceptable salts thereof. Claims 2 4, 5, 6 & 7, line 1: change "dithietan"

8. (amended) A process for the preparation of $\sqrt{7}$ (4-methoxy -7 β -(4-substituted methylene-1,3-diethietane-2-yl)carboxamido-3 -heterocyclic thiomethyl- Δ^3 -cephem-4-carboxylic acid represented by the [general] formula

to --dithietane-- (each occurrence, respectively).

 $R^{1} = S = S = CONH = CH_{2} - S - R^{3}$ $COOH = CH_{2} - S - R^{3}$

wherein R¹ represents a carboxyl group or the functional derivative residue thereof selected from the group consisting of a carboxylic acid lower alkyl ester residue, carboxylic acid aralkyl ester residue, carbamoyl, carbazolyl, and cyano groups; R² represents a

С

hydrogen atom, a lower alkyl group, a lower alkoxy group, R⁴S(O)_n group wherein R⁴ represents a lower alkyl group, and n represents 0, 1 or 2, a lower alkanoyl group, an aryl group an aroyl group, a carboxyl group or the functional derivative carboxyl group or the functional derivative carboxyl group consisting of a carboxylic acid lower ester residue, carboxylic acid aralkyl ester carboxylic acid lower ester residue, carboxylic acid aralkyl group, a sulfamoyl group, or a heterocyclic residue; and R³ represents a lower alkyl-substituted tetrazolyl group or a lower alkyl-substituted thiadiazolyl group [, R² and R³ have the same significance as in claim 1], which comprises reacting the 4-substituted methylene-1,3-dithietane-2 -carboxylic acid represented by the [general] formula

wherein R^1 and R^2 have the same significance as above, or the functional derivative thereof, with the 74-amino-76-methoxy-3--heterocyclic thiomethyl- \triangle^3 -cephem-4-carboxylic acid represented by the [general] formula

wherein R3 has the same significance as above.

preparation of a 9. (amended) A process for the/7%/methoxy-7%- (4-substituted methylene-1,3-dithietane-2-y1)carboxamido-3-heterocyclic thiomethyl- Δ^3 -cephem-4-carboxylic acid represented by the [general] formula

 $\begin{array}{c}
\text{OCH}_3 \\
\text{R}^2 =
\begin{array}{c}
\text{OCH}_3 \\
\text{ONH}
\end{array}$

represents a carboxyl group or the functional derivative from the group consisting of a carboxylic carboxylic acid aralkyl ester and and cyano groups; R2 represents a hydrogen atom, a lower alkyl group, a lower alkoxy group, represents a lower alkyl group and n represents or 2, a lower alkanoyl group, carboxyl group or the functional derivative the group consisting of a carboxylic acid lower/ester acid aralkvl ester lower alkenyl group, a sulfamoyl group, or a heterocyclic and R³ represents a lower alkylsubstituted tetrazolyl group or a lower alkyl-substituted thiadiazolyl group [, R^2 and R^3 have the same significance as in claim 1], which comprises reacting the 3-acetoxymethyl- (or 3-carbamoyl-oxymethyl-) 7(M-methox $\sqrt{-7}$)-(4-substituted methylene-1,3-dithietane-2-y1)carboxamido $-\Delta^3$ -cephem-4-carboxylic acid represented by the [general] formula

B 3

Þ

シ

wherein R^1 and R^2 have the same significance as above and R^6 represents an acetyl group or a carbamoyl group, with the heterocyclic thiol represented by the general formula

wherein R³ has the same significance as above.

10. (amended) A process for the preparation of $\sqrt{7}$ methoxy- 7β -(4-substituted methylene-), 3-dithietane-2-y1) carboxamido-3-heterocyclic thiomethyl- Δ 3-cephem-4-carboxylic acid represented by the [general] formula

wherein R² [and R³ have the same significance as in claim 1] represents a hydrogen atom, a lower alkyl group, a lower alkoxy group, R⁴S(O)_n group wherein R⁴ represents a lower alkyl group and n represents 0, 1 or 2, a lower alkanoyl group, an aryl group, an aroyl group, a carboxyl group or the functional derivative recidus thereof selected from the group consisting of a carboxylic acid alkyl lower ester residue, carboxylic acid aralkyl ester residue, carbamoyl, carbazologiand cyano groups, a lower alkenyl group, a sulfamoyl group, or a heterocyclic residue; and R³ represents a lower alkyl-substituted tetrazolyl group or a lower alkyl-substituted thiadiazolyl group, and R⁸ represents a hydrogen atom or a substituted or un-

7

substituted alkyl group, which comprises treating funder a basic condition] the 7%-methoxy-3-heterocyclic thiomethylcephalosporin derivative represented by the [general] formula

wherein R^2 , R^3 and R^8 have the same significance as above, with a

11. (amended) A 7 (3-Hydroxy-4-substituted isothiazol-5-yl) thioacetamido-7 (methoxy-3-heterocyclic thiomethyl-12-cephem-4-carboxylic acid represented by the [general] formula

HO R2 SCH₂CONH OCH₃ S CH₂-S-R

wherein R² [and R³ have the same significance as in claim 1.]

represents a hydrogen atom, a lower alkyl group, a lower alkoxy
group, R⁴S(O)
n group wherein R⁴ represents a lower alkyl group and
n represents 0, 1 or 2, a lower alkanoyl group, an aryl group, an
aroyl group, a carboxyl group or the functional derivative residue
thereof selected from the group consisting of a carboxylic acid
alkyl
lower /ester residue, carboxylic acid aralkyl ester residue,
carbamoyl, carboxer and cyano groups, a lower alkenyl group, a